

GenCore version 5.1.3
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OM protein - protein search, using sw model

Run on: November 30, 2002, 10:20:38 ; Search time 53.1921 Seconds
(without alignments)
2307.182 Million cell updates/sec

File: US-10-054-680-2

Perfect score: 4797
Sequence: 1 MAWLRLQLTSAFLHFLVLT.....LWLILYLPATLEAYCYIKGF 921

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 908470 seqs, 133250620 residues

Total number of hits satisfying chosen parameters: 908470

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database: A.Geneseq_101002:*

1:	/SIDS2/gcgdata/geneseq/genesqp-emb1/AA1980.DAT:*
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3:	/SIDS2/gcgdata/geneseq/genesqp-emb1/AA1982.DAT:*
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23:	/SIDS2/gcgdata/geneseq/genesqp-emb1/AA2002.DAT:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	4797	100.0	921	23	ABR3246
2	4784	99.7	927	23	AAW4745
3	4686	97.7	927	23	ABR3247
4	3425.5	71.4	970	23	AAE18291
5	3373.5	70.3	952	21	ABR41497
6	2143.5	44.7	609	22	ABR32633
7	2143.5	44.7	609	22	ABR18131
8	2143.5	44.7	609	22	AAW53461
9	2143.5	44.7	609	22	AAW13701
10	2143.5	44.7	609	22	AAW26102

11	2143.5	44.7	609	23	ABG35474	Human peptide enco
12	2130.5	44.4	950	22	ABR61721	Drosophila melanog
13	1607	33.5	394	22	ABG26781	Novel human diagno
14	589	12.5	120	22	AAO05893	Human polypeptide
15	579	12.1	539	21	AAV58044	Arabidopsis thailia
16	424	8.8	91	22	ABR27991	Human peptide #642
17	424	8.8	91	22	ABR3163	Peptide #669 enco
18	424	8.8	91	22	ABR18628	Protein #627 enco
19	424	8.8	91	22	AAW5959	Human brain expres
20	424	8.8	91	22	AAW6547	Human bone marrow
21	424	8.8	91	22	AAW14216	Peptide #650 enco
22	424	8.8	91	22	AAW26626	Peptide #663 enco
23	424	8.8	91	22	AAW01948	Peptide #630 enco
24	424	8.8	91	23	ABG35995	Human peptide enco
25	349	7.3	123	22	AAW23946	Rat EST encoded pr
26	239.5	5.0	618	22	AAW78712	Human protein Seq
27	234.5	4.9	603	23	AAE22088	Human novel lon ex
28	232.5	4.8	546	22	ABR12262	Human very large G
29	231.5	4.8	856	22	ABR60425	Drosophila melanog
30	227.5	4.7	658	22	ABR57820	Drosophila melanog
31	222	4.6	72	22	ABR15223	Human nervous syst
32	206	4.3	2071	22	AAE10949	Human protein sequ
33	206	4.3	2780	22	AAE10924	Human protein sequ
34	202	4.2	1615	23	ABR05663	Human signal trans
35	187	3.9	759	23	ABG61545	Human transporter
36	184	3.8	2777	22	AAE10925	Human monogenic au
37	181	3.8	1451	22	AAW95868	Human protein sequ
38	181	3.8	3530	22	AAW25586	Human protein sequ
39	179.5	3.7	3105	23	ABR80604	Human sbg101817CS
40	173.5	3.6	481	22	ABR58790	Drosophila melanog
41	165	3.4	353	23	AAE22090	Human novel lon ex
42	162.5	3.4	315	23	AAE22089	Human novel lon ex
43	157.5	3.3	480	22	ABG12623	Novel human diagno
44	157	3.3	168	20	AAV45263	Human secreted pro
45	151	3.1	42	22	ABR36549	Peptide #4055 enco

ALIGNMENTS

RESULT 1	
ABR3246	ABR3246 standard; Protein; 921 AA.
AC	ABR3246;
XX	
DT	21-AUG-2002 (first entry)
XX	
DE	Human transporter protein.
XX	
KW	Human; sodium/calcium exchanger; brain; heart; kidney; lung;
KW	spleen; testis; leukocyte; foetal brain; chromosome 1a.
OS	Homo sapiens.
XX	
PN	WO200233086-A2.
XX	
PD	25-APR-2002.
XX	
PF	17-OCT-2001; 2001WO-US32152.
XX	
PR	17-OCT-2000; 2000US-240836P.
XX	
PR	13-MAR-2001; 2001US-0804474.
XX	
PA	(PEKE) PE CORP NY.
XX	
PI	Merkulov GV, Ketchum KA, Shao W, Yan C, Di Francesco V;
XX	Beasley EM;
PI	WPI; 2002-479677/51.
DR	N-PSDB; ABRN83428, ABRN83429.
XX	
PT	Human transporter peptide related to sodium/calcium exchanger subfamily

PT for identifying modulators useful for treating a disease or condition
 PT mediated by human transporter protein -
 PS Claim 1; Fig 2; 200pp; English.

CC The present sequence is a human transporter protein, which is related to
 CC the sodium/calcium exchanger subfamily. Experimental data indicates
 CC expression of the transporter gene in humans in brain, heart, kidney,
 CC lung, spleen, testis, leukocyte and foetal brain. The gene of the
 CC transporter was mapped to chromosome 14 by ePCR.

XX Sequence 921 AA;

Query Match 100.0%; Score 4797; DB 23; Length 921;

Best Local Similarity 100.0%; Pred. No. 0;
 Matches 921; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MAMRLDPLTSAPLHFGVLFVFLNGLRAAGSGDVPSTGQNNESGSSDCKEGVIL 60
 DB 1 MAMRLDPLTSAPLHFGVLFVFLNGLRAAGSGDVPSTGQNNESGSSDCKEGVIL 60
 QY 61 PIWPEPNSLGDKTARVIVFVALIYMFVLSIADRFMAIEVITSQEREVTIKKPGE 120
 DB 61 PIWPEPNSLGDKTARVIVFVALIYMFVLSIADRFMAIEVITSQEREVTIKKPGE 120
 QY 121 TSTTIRVWNETVSNLTLMALGSSAPETILSLIEVCGHGFAGDLGPTIVGSAFNMFI 180
 DB 121 TSTTIRVWNETVSNLTLMALGSSAPETILSLIEVCGHGFAGDLGPTIVGSAFNMFI 180
 QY 181 IIGICVYIIPGGETRIKHLNRFVFIITAAWSTFAYITWLMIAVSPGVQVWEGILLTFE 240
 DB 181 IIGICVYIIPGGETRIKHLNRFVFIITAAWSTFAYITWLMIAVSPGVQVWEGILLTFE 240
 QY 241 FPVCVLLAMVADKRLFLFKYKHKYRTDKHRIIEETGHPKGIEMGKMMNSFLDGN 300
 DB 241 FPVCVLLAMVADKRLFLFKYKHKYRTDKHRIIEETGHPKGIEMGKMMNSFLDGN 300
 QY 301 LVPLEGKEVDSESRREMTIRILDKOKHPEKDLQLVEMANYALSHOOKSRAPYRIQATR 360
 DB 301 LVPLEGKEVDSESRREMTIRILDKOKHPEKDLQLVEMANYALSHOOKSRAPYRIQATR 360
 QY 361 MMTAGNLTLLKHAADQAKKASMSSEVHTDEPEDTSKYFPDPCSYQCIENCGAVLLTVR 420
 DB 361 MMTAGNLTLLKHAADQAKKASMSSEVHTDEPEDTSKYFPDPCSYQCIENCGAVLLTVR 420
 QY 421 KGGDMSTMYVDYKTEGOSANAGADYEFTGTVLAKPETOKESVGIIDDIIEEDBHF 480
 DB 421 KGGDMSTMYVDYKTEGOSANAGADYEFTGTVLAKPETOKESVGIIDDIIEEDBHF 480
 QY 481 FVRLSNVRIEEOPEEGMPAIFNSLPLPRAVLASPCVATVITLDDDHAGIFTECDTIH 540
 DB 481 FVRLSNVRIEEOPEEGMPAIFNSLPLPRAVLASPCVATVITLDDDHAGIFTECDTIH 540
 QY 541 VSESIGVMEVAVLTSGARGVIVPFRIVECTAGGGEDEFTDYGELFEKNDYVKIRV 600
 DB 541 VSESIGVMEVAVLTSGARGVIVPFRIVECTAGGGEDEFTDYGELFEKNDYVKIRV 600
 QY 601 KIYDEEYEROEENFFIALGEPKMERGISDVTDRKLFMEEBEAKRIEMGKPVVGEHPKL 660
 DB 601 KIYDEEYEROEENFFIALGEPKMERGISDVTDRKLFMEEBEAKRIEMGKPVVGEHPKL 660
 QY 661 EVIIEESYEFTKTVDKLIKKTNLALVYGTSHWRDQFMEAITVSAAGDEDESEGERLPS 720
 DB 661 EVIIEESYEFTKTVDKLIKKTNLALVYGTSHWRDQFMEAITVSAAGDEDESEGERLPS 720
 QY 721 CFEDVYMFELTYFKKVLFRACVPTEYCHGWACFAVSIILIGLTAIIGLASHFECTIGLK 780
 DB 721 CFEDVYMFELTYFKKVLFRACVPTEYCHGWACFAVSIILIGLTAIIGLASHFECTIGLK 780
 QY 781 DSVTAVFVAGTGVPTDFASKAALODVYADASIGNVTGSNAVNFLGIGLAVSAIY 840
 DB 781 DSVTAVFVAGTGVPTDFASKAALODVYADASIGNVTGSNAVNFLGIGLAVSAIY 840

QY 841 WALOGQEFHVSAGTLAFSVTLFTTFAFVCISVLLYRRRPHLGELGGRGCKLATWLEFV 900
 DB 841 WALOGQEFHVSAGTLAFSVTLFTTFAFVCISVLLYRRRPHLGELGGRGCKLATWLEFV 900
 QY 901 SLMLLYIFATLEAVCYIKGF 921
 DB 901 SLMLLYIFATLEAVCYIKGF 921

RESULT 2

AA047745
 ID AA047745 standard; Protein: 927 AA.

AC AA047745;
 XX

DT 25-FEB-2002 (first entry)
 XX

DE Human sodium(+)-calcium(2+) exchanger form 3 protein, HNCX3.

XX Human; Sodium(+)-Calcium(2+) exchanger form 3; HNCX3; chromosome 14;

XX cardiac failure; myocardial infarction; cardiac hypertrophy; arrhythmia;

XX myocarditis; pulmonary hypertension; cardiotoxicity; cardiac; Vaccine;

XX coronary heart disease; renal failure; ischaemic disorder;

XX Antiarrhythmic; Vasotropic; Hypotensive; cardiovascular disorder.

XX Homo sapiens.

PN W0200183744-A2.
 XX

PD 08-NOV-2001.
 XX

PE 30-APR-2001; 2001MO-EP04886.
 XX

PR 02-MAY-2000; 2000EP-0109080.
 XX

PA (MERE) MERCK PATENT GMBH.
 XX

PI Wilm C;
 XX

XX WPI: 2002-041493/05.
 DR

DR N-PSDB: ABA04756.
 XX

PT New polypeptide, useful as vaccines for inducing immune response
 PT against diseases such as myocardial infarction, arrhythmia, ischemic
 PT disorders, renal disorders in mammal -

XX Claim 1; Page 38-41; 41pp; English.

XX The present sequence is the protein sequence for human Sodium(+)-Calcium

XX (2+) exchanger form 3 (HNCX3). The HNCX3 gene maps to human chromosome

XX 14. HNCX3 and its coding sequence are useful for treating acute and

XX chronic cardiac failure of different aetiologies, myocardial infarction,

XX cardiac hypertrophy, arrhythmia, myocarditis, pulmonary hypertension,

XX cardiotoxicity (e.g. induced by chemotherapy), coronary heart disease,

XX acute and chronic renal failure, ischaemic disorders of skeletal muscle

XX and ischaemic brain disorders of different aetiologies.

XX Sequence 927 AA;

QY Query Match 99.7%; Score 4784; DB 23; Length 927;
 Best Local Similarity 99.4%; Pred. No. 0;
 Matches 921; Conservative 0; Mismatches 0; Indels 6; Gaps 1;

QY 1 MAMRLDPLTSAPLHFGVLFVFLNGLRAAGSGDVPSTGQNNESGSSDCKEGVIL 60
 DB 1 MAMRLDPLTSAPLHFGVLFVFLNGLRAAGSGDVPSTGQNNESGSSDCKEGVIL 60
 QY 61 PIWPEPNSLGDKTARVIVFVALIYMFVLSIADRFMAIEVITSQEREVTIKKPGE 120
 DB 61 PIWPEPNSLGDKTARVIVFVALIYMFVLSIADRFMAIEVITSQEREVTIKKPGE 120
 QY 121 TSTTIRVWNETVSNLTLMALGSSAPETILSLIEVCGHGFAGDLGPTIVGSAFNMFI 180
 DB 121 TSTTIRVWNETVSNLTLMALGSSAPETILSLIEVCGHGFAGDLGPTIVGSAFNMFI 180

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Db 121 TSTTIRVWNETVSNLTLMALGSSAPEILLSLIEVCGHFIAGDLPSTIVGSAFNMFI 180
Qy 181 IIGICVYVIPDGETRIKHLRVFETTAAMSTIFAYIMLWIIAIVSPGVQVWEGILLTFE 240
Db 181 IIGICVYVIPDGETRIKHLRVFETTAAMSTIFAYIMLWIIAIVSPGVQVWEGILLTFE 240
Qy 241 FPCVCLAMVADKRLLFYKMKKRYRTDKHRIIIEEGDHPKGIEMDGKMNSHFLDGN 300
Db 241 FPCVCLAMVADKRLLFYKMKKRYRTDKHRIIIEEGDHPKGIEMDGKMNSHFLDGN 300
Qy 301 LVPLEGGKVEDESRRERIRILKDLKOKHPEKDLQLEMANYYALSHOOKSRAFYRIQATR 360
Db 301 LVPLEGGKVEDESRRERIRILKDLKOKHPEKDLQLEMANYYALSHOOKSRAFYRIQATR 360
Qy 361 MMTGAGNLTAKHAABQAKKASSMSSEVHTDEPEDFISKVFEDCSYQCLENCAGVLLTVVR 420
Db 361 MMTGAGNLTAKHAABQAKKASSMSSEVHTDEPEDFISKVFEDCSYQCLENCAGVLLTVVR 420
Qy 421 KGDMSKTMVYDYKTEDGSANAGADYEFTEGTVLKPGETOKEFSGIIDDIFEEDEHF 480
Db 421 KGDMSKTMVYDYKTEDGSANAGADYEFTEGTVLKPGETOKEFSGIIDDIFEEDEHF 480
Qy 481 FVRLSNVRIEEBQPEEGMPALFNSLPLPRAVLASPCVATVITLDDHAGITFECDTIH 540
Db 481 FVRLSNVRIEEBQPEEGMPALFNSLPLPRAVLASPCVATVITLDDHAGITFECDTIH 540
Qy 541 VSESIGVMEVKVLRISGARGTVIPEPRIVEGTAKGGGEDEFTYGELEFKNDETYKTIHV 600
Db 541 VSESIGVMEVKVLRISGARGTVIPEPRIVEGTAKGGGEDEFTYGELEFKNDETYKTIHV 600
Qy 601 KIVDEEERENBNFIATGEPKMERGIS-----DVTDRKLTMEEEAKRIEMGKPYL 654
Db 601 KIVDEEERENBNFIATGEPKMERGIS-----DVTDRKLTMEEEAKRIEMGKPYL 654
Qy 655 GEHPRLEVIIEESYEFTKTVDKLIKLTALVVGTHSMRDQMEAITVSAAGDEDEDESG 714
Db 655 GEHPRLEVIIEESYEFTKTVDKLIKLTALVVGTHSMRDQMEAITVSAAGDEDEDESG 714
Qy 715 EERLPSCEVDYVHMLTIVYMKVLFACVPRTEYCHGWACRAVSLITIGMLAITGLASHHG 774
Db 715 EERLPSCEVDYVHMLTIVYMKVLFACVPRTEYCHGWACRAVSLITIGMLAITGLASHHG 774
Qy 775 CTIGLKDSVTAVFAFGTSVPDTPASKAALQDQVADASIGNVTGSNNANVFLGIGLAM 834
Db 775 CTIGLKDSVTAVFAFGTSVPDTPASKAALQDQVADASIGNVTGSNNANVFLGIGLAM 834
Qy 835 SVAALYMALQGEFHVSAAGTLAFSVTLFTFAFVCISVLLYRRPHLGGELGGPRGCKLA 894
Db 835 SVAALYMALQGEFHVSAAGTLAFSVTLFTFAFVCISVLLYRRPHLGGELGGPRGCKLA 894
Qy 895 TTWLFVSLMLLYIFATLEAYCYIKGF 921
Db 895 TTWLFVSLMLLYIFATLEAYCYIKGF 921
Qy 901 TTWLFVSLMLLYIFATLEAYCYIKGF 927
Db 901 TTWLFVSLMLLYIFATLEAYCYIKGF 927

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RESULT 3
ABB83247
ID ABB83247 standard; Protein: 927 AA.
AC ABB83247;
XX
XX 21-AUG-2002 (first entry)
XX
XX Human transporter protein-related protein, used in a homology alignment.
XX
XX Human; sodium/calcium exchanger; transporter; brain; heart; kidney; lung;
XX spleen; testis; leukocyte; foetal brain; chromosome 14.
XX
XX Unidentified.
XX
XX WO200233086-A2.
XX
XX 25-APR-2002.
PD

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XX 17-OCT-2001; 2001WO-US32152.
PF
XX 17-OCT-2000; 2000US-240836P.
PR
XX 13-MAR-2001; 2001US-0804474.
XX
XX (PEKE ) PE CORP NY.
XX
XX Merkulov GV, Ketchum KA, Shao W, Yan C, Di Francesco V;
XX Beasley EM;
XX WPI; 2002-479677/51.
XX
XX Human transporter peptide related to sodium/calcium exchanger subfamily
XX for identifying modulators useful for treating a disease or condition
XX mediated by human transporter protein
XX
XX Disclosure: Fig 2; 200pp; English.
XX
XX The present invention relates to a human transporter protein, which is
XX related to the sodium/calcium exchanger subfamily (ABB83246). In humans
XX Experimental data indicates expression of the transporter gene in
XX brain, heart, kidney, lung, spleen, testis, leukocyte and foetal
XX brain. The gene of the transporter was mapped to chromosome 14 by ePCR.
XX The present protein was used in a sequence alignment with the transporter
XX protein to illustrate the invention.
XX
XX Sequence 927 AA:
SQ

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Query Match 97.7%; Score 4686; DB 23; Length 927;
Best Local Similarity 96.8%; Pred. No. 0;
Matches 897; Conservative 14; Mismatches 10; Indels 6; Gaps 1;

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Qy 1 MAMRLQPLTSALFHLGTVFLVFLNGLAERAGSGDYVSTGONNESCSSDCKEGVIL 60
Db 1 MAMRLQPLTSALFHLGTVFLVFLNGLAERAGSLRDVPSAGONNESCSSDCKEGVIL 60
Qy 61 PIWYRPNPSLGKIRAVIYFVALIYMFVGSIIADRFMASTIEVTSQREVTIKKPNSE 120
Db 61 PIWYRPNPSLGKIRAVIYFVALIYMFVGSIIADRFMASTIEVTSQREVTIKKPNSE 120
Qy 121 TSTTIRVWNETVSNLTLMALGSSAPEILLSLIEVCGHFIAGDLPSTIVGSAFNMFI 180
Db 121 TSTTIRVWNETVSNLTLMALGSSAPEILLSLIEVCGHFIAGDLPSTIVGSAFNMFI 180
Qy 181 IIGICVYVIPDGETRIKHLRVFETTAAMSTIFAYIMLWIIAIVSPGVQVWEGILLTFE 240
Db 181 IIGICVYVIPDGETRIKHLRVFETTAAMSTIFAYIMLWIIAIVSPGVQVWEGILLTFE 240
Qy 241 FPCVCLAMVADKRLLFYKMKKRYRTDKHRIIIEEGDHPKGIEMDGKMNSHFLDGN 300
Db 241 FPCVCLAMVADKRLLFYKMKKRYRTDKHRIIIEEGDHPKGIEMDGKMNSHFLDGN 300
Qy 301 LVPLEGGKVEDESRRERIRILKDLKOKHPEKDLQLEMANYYALSHOOKSRAFYRIQATR 360
Db 301 LVPLEGGKVEDESRRERIRILKDLKOKHPEKDLQLEMANYYALSHOOKSRAFYRIQATR 360
Qy 361 MMTGAGNLTAKHAABQAKKASSMSSEVHTDEPEDFISKVFEDCSYQCLENCAGVLLTVVR 420
Db 361 MMTGAGNLTAKHAABQAKKASSMSSEVHTDEPEDFISKVFEDCSYQCLENCAGVLLTVVR 420
Qy 421 KGDMSKTMVYDYKTEDGSANAGADYEFTEGTVLKPGETOKEFSGIIDDIFEEDEHF 480
Db 421 KGDMSKTMVYDYKTEDGSANAGADYEFTEGTVLKPGETOKEFSGIIDDIFEEDEHF 480
Qy 481 FVRLSNVRIEEBQPEEGMPALFNSLPLPRAVLASPCVATVITLDDHAGITFECDTIH 540
Db 481 FVRLSNVRIEEBQPEEGMPALFNSLPLPRAVLASPCVATVITLDDHAGITFECDTIH 540
Qy 541 VSESIGVMEVKVLRISGARGTVIPEPRIVEGTAKGGGEDEFTYGELEFKNDETYKTIHV 600
Db 541 VSESIGVMEVKVLRISGARGTVIPEPRIVEGTAKGGGEDEFTYGELEFKNDETYKTIHV 600

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QY 844 OGOEHVSAGTLAFTVLTETFAVCISVLYRRRPHLGGELGPRGCKLATLWFLVSLM 903
 DB 893 NEOQFVSVSGTLAFTVLTETFAVCISVLYRRRPHLGGELGPRGCKLATLWFLVSLM 952
 QY 904 LYLIFATLEANCYIKGF 921
 DB 953 LYLIFATLEANCYIKGF 970

RESULT 5

AAB41497

ID AAB41497 standard; Protein; 952 AA.

AAB41497;

DT 08-FEB-2001 (first entry)

Human ORFX ORF1261 polypeptide sequence SEQ ID NO:2522.

Human: open reading frame; ORFX: detection; cytostatic; hepatotropic; vulnery; antiparietal; antiparkinsonian; neurotropic; neuroprotective; anticonvulsant; osteopathic; antidiabetic; immunosuppressant; cardiac; immunostimulant; thrombolytic; coagulant; vasotropic; antidiabetic; hypotensive; dermatological; immunosuppressive; antidiabetic; antilethal; antibacterial; antifungal; antineumatic; antihypertensive; antineumatic; gene therapy; cancer; proliferative disorder; hypertension; neurodegenerative disorder; osteoarthritis; graft vs host disease; cardiovascular disease; diabetes mellitus; hypothyroidism; SCID; AIDS; cholesterol ester storage; systemic lupus erythematosus; infection; severe combined immunodeficiency; malaria; autoimmune disorder; asthma; allergy; aplastic anaemia; nocturnal haemoglobinuria; burn; wound; bone damage; cartilage damage; antineumatic disease; coagulation; thrombosis; contraceptive.

Homo sapiens.

WO20058473-A2.

05-OCT-2000.

31-MAR-2000; 2000MO-US08621.

31-MAR-1999; 99US-0127607.

02-APR-1999; 99US-0127636.

05-APR-1999; 99US-0127728.

30-MAR-2000; 2000US-0540763.

(CURA-) CURAGEN CORP.

Shinkets RA, Leach M;

WPI. 2000-602362/57.

N-PSDB: AACT5706.

Claim 11: Page 1809-1812; 5507p; English.

AACT4446 to AACT7606 encode the proteins given in AAB40237 to AAB43397, which represent the human ORFX open reading frames 1 to 3161. The ORFX sequences have activities such as: cytostatic; hepatotropic; vulnery; antiparietal; antiparkinsonian; neurotropic; neuroprotective; anticonvulsant; osteopathic; antidiabetic; immunosuppressant; cardiac; immunostimulant; thrombolytic; coagulant; vasotropic; antidiabetic; hypotensive; dermatological; immunosuppressive; antineumatic; antibacterial; antifungal; antineumatic; antihypertensive; antineumatic. The sequences can be used for determining the presence of or predisposition to, or preventing or treating pathological conditions associated with an ORFX-associated disorder. The nucleic acids can be used to express ORFX proteins in gene therapy vectors. The proteins and nucleic acids may be used to treat cancers,

CC proliferative disorders, neurodegenerative disorders, osteoarthritis, CC graft vs host disease, cardiovascular disease, diabetes mellitus, CC erythematosus, severe combined immunodeficiency (SCID), AIDS, viral, CC bacterial or fungal infection, malaria, autoimmune disorders, asthma, CC allergies, aplastic anaemia, burns, wounds, bone and cartilage damage, CC nocturnal haemoglobinuria, antineumatic disease; to enhance CC coagulation; to inhibit thrombosis; and as a contraceptive.

Sequence 952 AA:

Query Match 70.3%; Score 3373.5; DB 21; Length 952;

Best local similarity 71.0%; Pred. No. 0; Matches 638; Conservative 124; Mismatches 104; Indels 33; Gaps 9;

QY 40 STGONNESCSDCKEGVILPIWYBNPSLQKARVYFVALYMFVLSIADREM 99
 DB 70 STG-----GCQSYRCOPGVLLPWEPDDPSLDKARAAYVYFVALYMFVLSIADREM 125
 QY 100 ASIEVITSOERETIKKPNGETSTTIRVNNETVSNLTLMALGSSAPEILLSIEVCGHG 159
 DB 126 AAEVITSKEKETITRKANGETSVGTRVWNETVSNLTLMALGSSAPEILLSIEVCGHN 185
 QY 160 FIAGDLPSTIVSAFNFNFIIGICVYVDPDETRIKHLRFVFTAAISFAYIWLVM 219
 DB 186 FOAGELGPGIYVSAFNFNFVAVNCIYIPAESKRIRKLRFPYATASIFAYIWLVL 245
 QY 220 ILAVFSPGVVWEGLLTFEPFVYLLAVNADRLLYKVMKKYRTDKHRIITETEG 279
 DB 246 ILAVFSPGVVWEGLLTFEPFVYLLAVNADRLLYKVMKKYRTDKHRIITETEG 305
 QY 280 DPKGIEIMCKMNSHFLQNTL-----VPLEGVEDSREMRIRLKLQKHPKEDLQ 334
 DB 306 DPKGIEIMCKMNSHFLQNTL-----VPLEGVEDSREMRIRLKLQKHPKEDLQ 364
 QY 335 LVEMANYVALSHOOKSRAFYRIQATRMKTCAGNLLKHAEOAKKASMSSEVTEDEDF 394
 DB 365 LVGIANVYALLHQQKSRAPRIQATRLMTGAGVLRHADAARRAARPAAGADEDE- DDG 423
 QY 395 ISKVFFDPCSYQCLNCGAVLLTVVRKGDMSKTMVYDYKTEGDSANAGADYFETGTV 454
 DB 424 ASRIFFEPSLYHCLNCGSVLLSVTCGCGGNSFTFYDRTEDGSAKAGSDYFSEGTLY 483
 QY 455 LKPGETOKERSVGIIDDDIFEEDHEFVRISNRIEEO-----PEGMPAIRNSLPLR 510
 DB 484 FKPGETOKERLIGIIDDIFEEDHEFVRISNRIEEO-----PEGMPAIRNSLPLR 534
 QY 511 AVLASPCVATVTLDDHAGIIFTECDTIHVSISIGMEVYKLVRTSGARCTVIVPRTVE 570
 DB 535 GRVAVPLATVTLDDHAGIIFTECDTIHVSISIGMEVYKLVRTSGARCTVIVPRTVE 594
 QY 571 GTAKGGGEDEFTYGELEFKNDETIVTRIKVIEDEEYERQENFTALGEPKMMERGISO 630
 DB 595 GTARGGVHVEDACGELEFEDDETMTLQVYKIVDEEYERQENFTALGEPKMMERGISA 654
 QY 631 VT-----DRKLMEEREARIAEMKRPVIGENCRLEVIIEESYDRKNTVDKLIKRTNLA 714
 DB 655 LLNQGDDGDKRLAEEERARIAEMKRPVIGENCRLEVIIEESYDRKNTVDKLIKRTNLA 744
 QY 685 LVVGTSMRDPMEALITVSAAGDEDEDESG--TEERLSPSCFDYVNHFTVMKLVLCVPP 742
 DB 715 LVVGTSMRDPMEALITVSAAGDEDEDESG--TEERLSPSCFDYVNHFTVMKLVLCVPP 773
 QY 743 TEYCHMACFVAVSILITGMILTAIGDASHFGCTIGLKDVSVAVFAVGTSPDTFASK 802
 DB 774 TEYCHMACFVAVSILITGMILTAIGDASHFGCTIGLKDVSVAVFAVGTSPDTFASK 833
 QY 803 AALADVVYDASIGNTGSAVNVFLGIGLAVVAALYALQOEFHVSAGTLAFTVTLR 862
 DB 834 VALADQCADASIGNTGSAVNVFLGIGLAVVAALYALQOEFHVSAGTLAFTVTLR 893
 QY 863 TIFAFVSVLTLRRRPHLGGELGPRGCKLATLWFLVSLMFLYIFATLEANCYIKGF 921

Db 894 TVFAFVGIALLVRRRRPHIGELGPGPKLATTALFLGLMLYLTFASLEAYCHINGF 952

RESULT 6
AB832633
ID AB832633 standard; Peptide: 609 AA.
XX
AC AB832633;
XX
DT 04-FEB-2002 (first entry)
XX
DE Peptide #139 encoded by human foetal liver single exon probe.
XX
KW Human; foetal liver; gene expression; single exon nucleic acid probe.
XX
OS Homo sapiens.
XX
PN MO200157277-A2.
XX
PD 09-AUG-2001.
XX
PR 30-JAN-2001; 2001WO-US00669.
XX
PR 04-FEB-2000; 2000US-0180312.
XX
PR 26-MAY-2000; 2000US-0207456.
XX
PR 30-JUN-2000; 2000US-0608408.
XX
PR 03-AUG-2000; 2000US-0632366.
XX
PR 21-SEP-2000; 2000US-0234687.
XX
PR 27-SEP-2000; 2000US-0236359.
XX
PR 04-OCT-2000; 2000GB-0024263.
XX
PA (MOLE-) MOLECULAR DYNAMICS INC.
XX
PI Penn SG, Hanzel DK, Chen W, Rank DR;
XX
XX WPI: 2001-483447/52.
XX
PT Human genome-derived single exon nucleic acid probes useful for
XX analyzing gene expression in human fetal liver -
XX
PS Claim 27: SEQ ID NO 25268; 639pp + sequence listing; English.
XX
CC The invention relates to a single exon nucleic acid probe for
XX measuring human gene expression in a sample derived from human foetal
XX liver. The single exon nucleic acid probes may be used for predicting,
XX measuring and displaying gene expression in samples derived from human
XX foetal liver. The present sequence is a peptide encoded by a single exon
XX nucleic acid probe of the invention.
XX
CC Note: The sequence data for this patent did not form part of the
XX printed specification, but was obtained in electronic format directly
XX from WIPO at ftp.wipo.int/pub/published_pct_sequences.
XX
SQ Sequence 609 AA;
XX
Query Match 44.7%; Score 2143.5; DB 22; Length 609;
Best Local Similarity 69.3%; Pred. No. 6.9e-200;
Matches 420; Conservative 76; Mismatches 89; Indels 21; Caps 8;
XX
QY 1 MAMRLQPLTSGAFHFGVTFVLF--LNGLRAGGSGDVSTGONNESCSCGSDCKREGV 58
DB 11 MRLSLSPTEFSGFHLVTVSLFSHDVIAETEMEGETGE---CTGSYCKKGV 66
XX
QY 59 ILPTWYENPSPISGDKIARVIYFVALIYMFGLVSIADRFMASTEVITSOBREVITIKPN 118
DB 67 ILPTWYENPSPISGDKIARVIYFVALIYMFGLVSIADRFMASTEVITSOBREVITIKPN 126
XX
QY 119 GETSTTIRWNETVSNLTLMALGSSAPEILLSEVCGHGFAGDLPSTIVGSAFNM 178
DB 127 GETTKTVRIWNETVSNLTLMALGSSAPEILLSEVCGHGFAGDLPSTIVGSAFNM 186
XX
QY 179 FIITGICVYVPDGETRIKRLRVFTITAMSFATYIMLITLAVFSGVQVWEGLLTL 238
DB 187 FIITACVYVVPDGETRIKRLRVFTITAMSFATYIMLITLAVFSGVQVWEGLLTL 246
XX

QY 239 FFFPVYLLAVNADKRLLEFYVMHKKYRTDKHRIIETEGDHPKG---IMDGKMNSH 295
DB 247 FFFPICVYFAMVADRRLLEFYKYYKRYRAGKQRMIIIEHEDGRSSKTEIMDGKVNSH 306
XX
QY 296 ---FLDGLNVPLEGKVD---ESRREMIKRLKDKQKHPKDDQVLEMANYYALSIQQ 348
DB 307 VENFLDGLNVPLEGKVD---ESRREMIKRLKDKQKHPKDDQVLEMANYYALSIQQ 365
XX
QY 349 KSRAFYRIQATRMNTGACNIIKHAADQAKASSSEVHTDEPE-DFISKYFDPSCSYQC 407
DB 366 KSRAFYRIQATRMNTGACNIIKHAADQAKASSSEVHTDEPE-DFISKYFDPSCSYQC 425
XX
QY 408 LENCQAVLLTVYRKGDMSKTYVDYKTEDGSANAGADYEFTGTYVLLKPGETKESVGC 467
DB 426 LENCQAVLLTVYRKGDMSKTYVDYKTEDGSANAGADYEFTGTYVLLKPGETKESVGC 485
XX
QY 468 IIDDIFEDDEHFFVRLSNVRIEEOPEEGPAPLPLPRAVLASPCVATYTIILDD 527
DB 486 IIDDIFEDDEHFFVRLSNVRIEEOPEEGPAPLPLPRAVLASPCVATYTIILDD 542
XX
QY 528 HAGIFTEPCDTIHYSESIGVMEVVLRTSGARGVYVPEFTVEGTAGGGEFEDTYGEL 587
DB 543 HAGIFTEPCDTIHYSESIGVMEVVLRTSGARGVYVPEFTVEGTAGGGEFEDTYGEL 602
XX
QY 588 EFKNDE 593
DB 603 EFKNDE 608
XX
RESULT 7
AB818131
ID AB818131 standard; Protein: 609 AA.
XX
AC AB818131;
XX
DT 23-JAN-2002 (first entry)
XX
DE Protein #130 encoded by probe for measuring heart cell gene expression.
XX
KW Human; gene expression; heart; microarray; vascular system;
KW cardiovascular disease; hypertension; cardiac arrhythmia;
KW congenital heart disease.
XX
OS Homo sapiens.
XX
PN MO200157274-A2.
XX
PD 09-AUG-2001.
XX
PE 30-JAN-2001; 2001WO-US00666.
XX
PR 04-FEB-2000; 2000US-0180312.
XX
PR 26-MAY-2000; 2000US-0207456.
XX
PR 30-JUN-2000; 2000US-0608408.
XX
PR 03-AUG-2000; 2000US-0632366.
XX
PR 21-SEP-2000; 2000US-0234687.
XX
PR 27-SEP-2000; 2000US-0236359.
XX
PR 04-OCT-2000; 2000GB-0024263.
XX
PA (MOLE-) MOLECULAR DYNAMICS INC.
XX
PI Penn SG, Hanzel DK, Chen W, Rank DR;
XX
XX WPI: 2001-488899/53.
XX
PT Single exon nucleic acid probes for analyzing gene expression in human
XX hearts -
XX
PS Claim 15: SEQ ID NO 19901; 530pp; English.
XX
CC The present invention relates to single exon nucleic acid probes for
XX measuring human gene expression in a sample derived from human heart (see

CC ABA21535-ABA41305). The present sequence is a protein encoded by one such
 CC probe. The probes may be used for predicting, measuring and displaying
 CC gene expression in samples derived from the human heart via microarrays.
 CC By measuring gene expression, the probes are useful for predicting,
 CC diagnosing, grading, staging, monitoring and prognosing diseases of the
 CC human heart and vascular system e.g. cardiovascular disease,
 CC hypertension, cardiac arrhythmias and congenital heart disease.
 CC Note: The sequence data for this patent did not form part of the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pcl_sequences.

xx Sequence 609 AA:

Query Match 44.7%; Score 2143.5; DB 22; Length 609;

Best Local Similarity 69.3%; Pred. No. 6.9e-200;

Matches 420; Conservative 76; Mismatches 89; Indels 21; Gaps 8;

QY 1 MAMRLQPLTSAFLHGLVTVLFL--LNGLRAGAGSGDVPSTGQNNESGSSDCKEY 58
 DB 11 MRLSLSPFMSGPHLVVSLFSDHVDVIAETEMEGENETGE---CTGSYCKKV 66
 QY 59 ILPIWPNPNSLGDKTARIVYFVALIYMLFVSIADRFMAISIEVTSQEREVTIKKP 118
 DB 67 ILPIWPNPNSLGDKTARIVYFVALIYMLFVSIADRFMAISIEVTSQEREVTIKKP 126
 QY 119 GETSTTIRVWNETVSNLTLMALGSSAPETLLIEVCGHGFACDLSPTIYGAFAFM 178
 DB 127 GETTCTTIRVWNETVSNLTLMALGSSAPETLLIEVCGHGFACDLSPTIYGAFAFM 186
 QY 179 FTIIGICVYVDPGETRRIKHLRFVFTTAAWSIFAYIWMILAVSPGVQVWEGLLTL 238
 DB 187 FTIIGICVYVDPGETRRIKHLRFVFTTAAWSIFAYIWMILAVSPGVQVWEGLLTL 246
 QY 239 FFFPVCVLLAVADKRLFLFYKYMHHKRYRDKHNGIITETGDKPKG---TEMGKMNNSH 295
 DB 247 FFFPVCVLLAVADKRLFLFYKYMHHKRYRDKHNGIITETGDKPKG---TEMGKMNNSH 306
 QY 296 ---FLDGNLVPLEKGEVD---ESRREMIIRLKDLOKHPKDLQVEMANYALSHQ 348
 DB 307 VENFDGALV-LEVDERDODDEARREMAKILKELKOKHPDKIEBOLIELANTOVLSDQ 365
 QY 349 KSRAFYRIQATRMGTAGNLIKRAAQAQKASMSSEVHTDEPE-DFISKVFDPSCSYOC 407
 DB 366 KSRAFYRIQATRMGTAGNLIKRAAQAQKASMSSEVHTDEPE-DFISKVFDPSCSYOC 425
 QY 408 LENCGAVLLTVKKGDMKTMVDYKTEDEGSANAGADYEFTGTVLAKRGETQKEFSVG 467
 DB 426 LENCGAVLLTVKKGDMKTMVDYKTEDEGSANAGADYEFTGTVLAKRGETQKEFSVG 485
 QY 468 IIDDIDFEEDHFFVLSNVRIEEOPEBGMPPAIFNSLPLPRAVLASPCVALVITLDD 527
 DB 486 IIDDIDFEEDHFFVLSNVRIEEOPEBGMPPAIFNSLPLPRAVLASPCVALVITLDD 542
 QY 528 HAGIFTECDTIHVSISIGMEVYKLTSGARGTVIYFRTVEGTAKGGEDEEDTYGEL 587
 DB 543 HAGIFTECDTIHVSISIGMEVYKLTSGARGTVIYFRTVEGTAKGGEDEEDTYGEL 602
 QY 588 EFKNDE 593
 DB 603 EFKNDE 608
 RESULT 8
 AAM53461
 ID AAM53461 standard; Protein; 609 AA.
 AC AAM53461;
 XX 05-NOV-2001 (first entry)
 DT Human brain expressed single exon probe encoded protein seq ID NO: 25566.
 DE Human brain expressed single exon probe encoded protein seq ID NO: 25566.
 XX Human; brain expressed exon; gene expression analysis; probe;

KW microarray; Alzheimer's disease; multiple sclerosis; schizophrenia;
 KW epilepsy; cancer.

OS Homo sapiens.

PN W0200157275-A2.

PD 09-AUG-2001.

PF 30-JAN-2001; 2001WO-US00667.

PR 04-FEB-2000; 2000US-0180312.

PR 26-MAY-2000; 2000US-0207456.

PR 30-JUN-2000; 2000US-0608408.

PR 03-AUG-2000; 2000US-0632366.

PR 21-SEP-2000; 2000US-0234687.

PR 27-SEP-2000; 2000US-0236359.

PR 04-OCT-2000; 2000GB-0024263.

XX (MOLE-) MOLECULAR DYNAMICS INC.

XX Penn SG, Hanzel DK, Chen W, Rank DR;

XX WPI; 2001-483446/52.

XX Single exon nucleic acid probes for analyzing gene expression in human

XX brains -

XX Example 4; SEQ ID NO: 25566; 650pp + sequence listing; English.

XX The present invention provides a number of single exon nucleic acid

XX probes which are derived from genomic sequences expressed in the human

XX brain. They can be used to measure gene expression in brain cell samples,

XX which may enable the diagnosis and improved treatment of nervous system

XX diseases such as Alzheimer's disease, multiple sclerosis, schizophrenia,

XX epilepsy and cancers. The present sequence is a protein encoded by one of

XX the probes of the invention.

SEQ Sequence 609 AA:

Query Match 44.7%; Score 2143.5; DB 22; Length 609;

Best Local Similarity 69.3%; Pred. No. 6.9e-200;

Matches 420; Conservative 76; Mismatches 89; Indels 21; Gaps 8;

QY 1 MAMRLQPLTSAFLHGLVTVLFL--LNGLRAGAGSGDVPSTGQNNESGSSDCKEY 58
 DB 11 MRLSLSPFMSGPHLVVSLFSDHVDVIAETEMEGENETGE---CTGSYCKKV 66
 QY 59 ILPIWPNPNSLGDKTARIVYFVALIYMLFVSIADRFMAISIEVTSQEREVTIKKP 118
 DB 67 ILPIWPNPNSLGDKTARIVYFVALIYMLFVSIADRFMAISIEVTSQEREVTIKKP 126
 QY 119 GETSTTIRVWNETVSNLTLMALGSSAPETLLIEVCGHGFACDLSPTIYGAFAFM 178
 DB 127 GETTCTTIRVWNETVSNLTLMALGSSAPETLLIEVCGHGFACDLSPTIYGAFAFM 186
 QY 179 FTIIGICVYVDPGETRRIKHLRFVFTTAAWSIFAYIWMILAVSPGVQVWEGLLTL 238
 DB 187 FTIIGICVYVDPGETRRIKHLRFVFTTAAWSIFAYIWMILAVSPGVQVWEGLLTL 246
 QY 239 FFFPVCVLLAVADKRLFLFYKYMHHKRYRDKHNGIITETGDKPKG---TEMGKMNNSH 295
 DB 247 FFFPVCVLLAVADKRLFLFYKYMHHKRYRDKHNGIITETGDKPKG---TEMGKMNNSH 306
 QY 296 ---FLDGNLVPLEKGEVD---ESRREMIIRLKDLOKHPKDLQVEMANYALSHQ 348
 DB 307 VENFDGALV-LEVDERDODDEARREMAKILKELKOKHPDKIEBOLIELANTOVLSDQ 365
 QY 349 KSRAFYRIQATRMGTAGNLIKRAAQAQKASMSSEVHTDEPE-DFISKVFDPSCSYOC 407
 DB 366 KSRAFYRIQATRMGTAGNLIKRAAQAQKASMSSEVHTDEPE-DFISKVFDPSCSYOC 425
 QY 408 LENCGAVLLTVKKGDMKTMVDYKTEDEGSANAGADYEFTGTVLAKRGETQKEFSVG 467

Db		426	LKMGCVALLTIIRGGDITNTVFDRTEDGTANAGSDYFTGCTVVKFGDQKEIRVG	485
Qy		468	IIDDDIFEEDEHFFVRLSNRRIEEOPEEGMPAIFNSLPLPRAVLASPCVAATYILDD	527
Db		486	IIDDIFEEDEENFLVHLSNVKSSEASEDGILEANHVS--TLACLGSPSTAFVTIPDD	542
Qy		528	HAGIIFEEDCDTHVSESIGMEYKVLTSGARCTVIYPFVTGCTAKGGDEPDYTIGEL	587
Db		543	HAGIIFEFEVPVTHVSESIGIMEYKVLTSGARNVIVPYKTICTANGGGEDEPDITGEL	602
Qy		588	EFKNDE 593 :	
Db		603	EFGNDE 608	
RESULT 9				
AAM13701				
ID		AAM13701	standard; Protein; 609 AA.	
XX				
AC		AAM13701;		
XX				
XT		12-OCT-2001	(first entry)	
XX				
DE		Pepide #135 encoded by probe for measuring cervical gene expression.		
XX		Probe; human; microarray; gene expression; cervical epithelial cell;		
KM		cervical cancer.		
XX				
OS		Homo sapiens.		
PN		WO200157278-A2.		
XX				
PD		09-AUG-2001.		
XX				
PF		30-JAN-2001; 2001MO-US006070.		
XX				
PR		04-FEB-2000; 2000US-0180312.		
PR		26-MAY-2000; 2000US-0207456.		
PR		30-JUN-2000; 2000US-0608408.		
PR		03-AUG-2000; 2000US-0632366.		
PR		21-SEP-2000; 2000US-0234687.		
PR		27-SEP-2000; 2000US-0236359.		
PR		04-OCT-2000; 2000SB-0024263.		
PA		(MOLE-) MOLECULAR DYNAMICS INC.		
XX				
PI		Penn SG, Hanzel DK, Chen W, Rank DR;		
XX				
DR		WPT; 2001-488901/53.		
XX				
PT		Human genome-derived single exon nucleic acid probes useful for		
XX		analyzing gene expression in human cervical epithelial cells -		
PS		Claim 27; SEQ ID No 18527; 487bp; English.		
XX				
CC		The present invention relates to human single exon nucleic acid probes		
CC		(SENPs: see AAI10068-AA18459). The present sequence is a peptide encoded		
CC		by one such probe. The SENPs are derived from human HeLa cells. The SENPs		
CC		can be used to produce a single exon microarray, which can be used for		
CC		measuring human gene expression in a sample derived from human cervical		
CC		epithelial cells. By measuring gene expression, the probes are therefore		
CC		useful in grading and/or staging of diseases of the cervix, notably		
CC		cervical cancer.		
CC		Note: The sequence data for this patent did not form part of the printed		
CC		specification, but was obtained in electronic format directly from WIPO		
CC		at ftp.wipo.int/pub/published_pct_sequences.		
XX				
SQ		Sequence 609 AA:		
Query Match		44.7%; Score 2143.5; DB 22; Length 609;		
Best Local Similarity		69.3%; Pred. No. 6.9e-200;		
Matches 420; Conservative 76; Mismatches 89; Indels 21; Gaps 8				

QY	1	MAHLAIGLITSAFLHFGVLTVEF--LNGLRAEAGSGDVPSYGNONNCSGSGSDCKEY	58
Db	11	MRRLSLPTFSNGRHLVTWSLSLSDHDAIAETEMGEENETGE---CTGYTCKKCV	66
QY	59	ILPIVPPNPSPISGKIAIAVIVYFVALIYMFGLVSIIDRFMASIEVTSOEREVTIKKN	118
Db	67	ILPIWPPDPSPGDKIAIAATVYFVAAMYMFGLSIIDRFMSSIIVITSGEKTITKKNN	126
QY	119	GETSTTTIRVNNENVSNLTLMALGSSAPEILLSLIEVCGGFIAGDLGPSTIYGSAAFM	178
Db	127	GETTETVRIANNENVSNLTLMALGSSAPEILLSVIEVCGHNFAGDLGESTIYGSAAFM	186
QY	179	FLIIICGVYVPDGETRIKHLRFVFTTAAMSIFAYTMLMIILAVSPGVQVWEGLLT	238
Db	187	FLIILACVYVPDGETRIKHLRFVFTTAAMSIFAYTMLIILSVISPGVYEWEGLLTF	246
QY	239	FEFPVCVLLAVYADKRLLFVYKMKHYKRTDKHRCIILETEGDHPKG--IEMDGKMMNSH	295
Db	247	FEFPICVYFAWVADBRLLFVYKYVKRYRACKQRMILIEHGDPRSCKTEIEMDGKVNSH	306
QY	296	---FLDGLVLYELEGKVD---EERREMIRLDLQKHREKRLDQVLEKANTYALSHQ	348
Db	307	VENFLDGALV-LVDYERDQDDEEARREMARILKELQKHDKIEIDLIELANQVLSQO	365
QY	349	KSAFYRIQATRRMTGAGNIIKTKAAQAKKASMSSEVHTDEDE-DFISKVFPDPCSLOC	407
Db	366	KSAFYRIQATRLMTGAGNIIKRIAAQAKKASMSHEVNTENVENDPYSKIFPEQGTYC	425
QY	408	LENGCAVLLLVYRKGGDKMTMYVDYTEDGSSANAGADVEFTGETVLYLPGETQKEFSVG	467
Db	426	LENGCTVALTLIRRGDLTLNTEVFDFRTEDGTANAGSDVEFTGETVLYFKRGDTQKEIRVG	485
QY	468	IIDDIFEDDEHFFVYRLSNVRIIEEQDEEGMPRAIFNSLPLPRAYLASCVATVITLDD	527
Db	486	IIDDIFEDDENFLVHLSNNKVSSEASESDGILLANHVS--TLACIGSPSTAVITLITDD	542
QY	528	HAGITFECDTIHSESIGVMEVVLRTSGARGVIVPFTFVEGTAKGGEDFEDTYGEL	587
Db	543	HAGITFEPEPTIHSSESIGIMEVVLRTSGARGVIVPYKTIGTARGGGDEPDTGSEL	602
QY	588	EFRKND E 593	
Db	603	EFRKND E 608	
RESULT 10			
AAW26102	ID	AAW26102 standard; Protein; 609 AA.	
AC	XX	AAW26102;	
XX	XX	17-OCT-2001 (first entry)	
DE	XX	Peptide #139 encoded by probe for measuring placental gene expression	
KN	XX	Probe; microarray; human; placenta; antenatal diagnosis;	
KW	XX	genetic disorder.	
OS	XX	Homo sapiens.	
PN	XX	MO200157272-A2.	
PD	XX	09-AUG-2001.	
PF	XX	30-JAN-2001; 2001WO-0500663.	
PR	XX	04-FEB-2000; 2000US-0180312.	
PR	XX	26-MAY-2000; 2000US-0207456.	
PR	XX	30-JUN-2000; 2000US-0608408.	
PR	XX	03-AUG-2000; 2000US-0632366.	
PR	XX	21-SEP-2000; 2000US-0234687.	
PR	XX	27-SEP-2000; 2000US-0236359.	

PR 04-OCT-2000; 2000GB-0024263.
 XX (MOLE-) MOLECULAR DYNAMICS INC.
 XX Penn SG, Hanzel DK, Chen W, Rank DR;
 PI WPI; 2001-488897/53.
 XX
 XX WPI; 2001-488897/53.
 PT Human genome-derived single exon nucleic acid probes useful for
 PT analyzing gene expression in human placenta -
 XX
 XX Claim 27; SEQ ID No 26371; 654bp; English.
 PS
 CC The present invention relates to single exon nucleic acid probes (SENPs;
 CC see A131315-A157546). The present sequence is a peptide encoded by one
 CC such probe. The probes are useful for producing a microarray for
 CC predicting, measuring and displaying gene expression in samples derived
 CC from human placenta. The probes are useful for antenatal diagnosis of
 CC human genetic disorders.
 CC
 SO Sequence 609 AA:
 Query Match 44.7%; Score 2143.5; DB 22; Length 609;
 Best Local Similarity 69.3%; Pred. No. 6.9e-200;
 Matches 420; Conservative 76; Mismatches 89; Indels 21; Gaps 8;
 QY 1 MMLRLQPLTSAFLHFGVTLFVLF--LNLRLAAGSGSDVPSTGQNNESGSSDCKEY 58
 DB 11 MRLSLSPFSGFHLVTVSLFSLVHDVIAETEMEGSGENEGE---CTGSGYCKKG 66
 QY 59 ILPIWYENPSPGLDKIARIYVYFVALIYMLGSIADRFMASIEVITSOERVTIKRP 118
 DB 67 ILPIWYENPSPGLDKIARIYVYFVALIYMLGSIADRFMASIEVITSOERVTIKRP 126
 QY 119 GETSTTIRVWNETVSNLTALMGSSAPILLSLFVCGHFIAGDLGPSTVIGSAFNM 178
 DB 127 GETTCTVIMNETVSNLTALMGSSAPILLSLFVCGHFIAGDLGPSTVIGSAFNM 186
 QY 179 FTIIGCVVIVDPGETRKRKHLRFVITAASTFAITIMLAVSPGVQVWEGILL 238
 DB 187 FTIIGCVVIVDPGETRKRKHLRFVITAASTFAITIMLAVSPGVQVWEGILL 246
 QY 239 FFFPVCLVLAADKRLFYKYHKKYKRDHKGIIIEGDKPKG---IEMGKMNNSH 295
 DB 247 FFFPVCLVLAADKRLFYKYHKKYKRDHKGIIIEGDKPKG---IEMGKMNNSH 306
 QY 296 ---FLDGNLVPLEGKEVD---ESRREMTIRILDKOKHPEKDDQJLVEMANYALSHQ 348
 DB 307 VENFLDGNLVPLEGKEVD---ESRREMTIRILDKOKHPEKDDQJLVEMANYALSHQ 365
 QY 349 KSRATFRIDATNMGTGAGILKKHAQAOKKASSKSEVHTDEPE-DTISKVFPDPCSYOC 407
 DB 366 KSRATFRIDATNMGTGAGILKKHAQAOKKASSKSEVHTDEPE-DTISKVFPDPCSYOC 425
 QY 408 LENCAGVLLTVVRKGGDKMTYVDKTTEDGSANAGADFEPTGVLLRPGEOKEFSYG 467
 DB 426 LENCAGVLLTVVRKGGDKMTYVDKTTEDGSANAGADFEPTGVLLRPGEOKEFSYG 485
 QY 468 IIDDIFEEDEHFEVRLSNVRIEEOPEEGMPAIFNSLPLPRAVLASPCVATVITLDD 527
 DB 486 IIDDIFEEDEHFEVRLSNVRIEEOPEEGMPAIFNSLPLPRAVLASPCVATVITLDD 542
 QY 528 HAGIFFECDTHVSEISIGVMEKVVLRITSGAKTIVVPRVTVGAKKGGGEDEFTYGL 567
 DB 543 HAGIFFECDTHVSEISIGVMEKVVLRITSGAKTIVVPRVTVGAKKGGGEDEFTYGL 602
 QY 588 EFKNDE 593
 DB 603 EFKNDE 608

ID ABG35474 standard; Peptide; 609 AA.
 XX
 AC ABG35474;
 XX
 DT 19-AUG-2002 (first entry)
 XX
 DE Human peptide encoded by genome-derived single exon probe SEQ ID 25139.
 XX
 KW Human; single exon probe; asthma; lung cancer; COPD; ILD;
 KW chronic obstructive pulmonary disease; interstitial lung disease;
 KW familial idiopathic pulmonary fibrosis; neurofibromatosis;
 KW tuberous sclerosis; Gaucher's disease; Niemann-Pick disease;
 KW Hereditary spherocytosis; sarcoidosis; pulmonary haemorrhage;
 KW pulmonary histiocytosis; lymphangioleiomyomatosis; Kargener syndrome;
 KW pulmonary alveolar proteinosis; fibrocystic pulmonary dysplasia;
 KW primary ciliary dyskinesia; pulmonary hypertension;
 KW hyaline membrane disease.
 KW
 OS Homo sapiens.
 XX
 PN WO200186003-A2.
 XX
 PD 15-NOV-2001.
 XX
 PF 30-JAN-2001; 2001WO-US00665.
 XX
 PR 04-FEB-2000; 2000US-180312P.
 PR 26-MAY-2000; 2000US-207456P.
 PR 30-JUN-2000; 2000US-0608408.
 PR 03-AUG-2000; 2000US-0632366.
 PR 21-SEP-2000; 2000US-234687P.
 PR 27-SEP-2000; 2000US-236359P.
 PR 04-OCT-2000; 2000GB-0024263.
 XX
 PA (MOLE-) MOLECULAR DYNAMICS INC.
 PI Penn SG, Hanzel DK, Chen W, Rank DR;
 DR WPI; 2002-114183/15.
 XX
 PT Spatially-addressable set of single exon nucleic acid probes, used to
 PT measure gene expression in human lung samples -
 XX
 PS Claim 27; SEQ ID No 25139; 634bp; English.
 XX
 CC The invention relates to a spatially-addressable set of single exon
 CC nucleic acid probes for measuring gene expression in a sample derived
 CC from human lung comprising single exon nucleic acid probes having one of
 CC 12614 nucleic acid sequences mentioned in the specification, or their
 CC complements or the 12614 open reading frames derived from the 12614
 CC probes. Also included are a microarray comprising the novel set of
 CC probes; the novel set of probes which hybridize at high stringency to a
 CC nucleic acid expressed in the human lung; measuring gene expression in a
 CC sample derived from human lung, comprising (a) contacting the array with
 CC a collection of detectably labeled nucleic acids derived from human lung
 CC mRNA, and (b) measuring the label detectably bound to each probe of
 CC the array; identifying exons in a eukaryotic genome, comprising
 CC (a) algorithmically predicting at least one exon from genomic sequences
 CC of the eukaryote; and (b) detecting specific hybridisation of detectably
 CC labeled nucleic acids from eukaryotic lung mRNA, to a single exon probe,
 CC having a fragment identical to the predicted exon, the probe is included
 CC in the above mentioned microarray, assigning exons to a single gene,
 CC comprising (a) identifying exons from genomic sequence by the method
 CC above and (b) measuring the expression of each of the exons in several
 CC tissues and/or cell types using hybridisation to a single exon
 CC microarrays having a probe with the exon, where a common pattern of
 CC expression of the exons in the tissues and/or cell types indicates that
 CC the exons should be assigned to a single gene; a peptide comprising one
 CC of 12011 sequences, mentioned in the specification, or encoded by the
 CC probes/open reading frames (ORF). The probes are used for gene
 CC expression analysis, and for identifying exons in a gene, particularly
 CC using human lung derived mRNA and for the study of lung diseases
 CC such as asthma, lung cancer, chronic obstructive pulmonary disease

RESULT 11
 ABG35474

CC (COPD), interstitial lung disease (ILD), familial idiopathic pulmonary
 CC fibrosis, neurofibromatosis, tuberous sclerosis, Gaucher's disease,
 CC Mleman-Pick disease, Hermansky-Pudlak syndrome, sarcoidosis, pulmonary
 CC hemangioendothelioma, pulmonary histiocytosis, lymphangioleiomyomatosis,
 CC pulmonary alveolar proteinosis, Kargener syndrome, fibrocystic
 CC pulmonary dysplasia, primary ciliary dyskinesia, pulmonary hypertension
 CC and hyaline membrane disease. The present sequence is a peptide/protein
 CC encoded by a single exon probe of the invention.
 CC Note: The sequence data for this patent did not form part
 CC of the printed specification, but was obtained in electronic
 CC format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences.

XX Sequence 609 AA;

Query Match 44.7%; Score 2143.5; DB 23; Length 609;
 Best Local Similarity 69.3%; Pred. No. 6.9e-200;

Matches 420; Conservative 76; Mismatches 89; Indels 21; Gaps 8;

QY 1 MAMRLQPLTSAPLHFGVLFVLF-LNGLRAGSGSDVPTGONNESCSSGSDCKEGV 58
 DB 11 MRRLSLSPFTSMGPHLLVTVSLFSDHVAIETEMEGEGTEGE---CTGTYCKKGV 66
 QY 59 ILPLTWENPSLQKIRARIVYFVALIYMLGVSIIADREMAIEVITTSQREVTIKPN 118
 DB 67 ILPLTWENPSLQKIRARIVYFVALIYMLGVSIIADREMAIEVITTSQREVTIKPN 126
 QY 119 GETSTTIRVWNETVSNLTLMALGSSAPETLSIEVCGHGFITGDIQSPSTIVSAFNM 178
 DB 127 GETTKTVIRINETVSNLTLMALGSSAPETLSIEVCGHGFITGDIQSPSTIVSAFNM 186
 QY 179 FIIGICVYVDPDETRKIKHLRVFTTAWSIFAYIMLYMILAVFSPGVQVVEGLTTL 238
 DB 187 FIILACVYVDPDETRKIKHLRVFTTAWSIFAYIMLYMILAVFSPGVQVVEGLTTL 246
 QY 239 FEFVVCVLAVNADKRLFFYMKKRTDKHKGIIETESDHRG---IEMDKMNSH 295
 DB 247 FEFVVCVLAVNADKRLFFYMKKRTDKHKGIIETESDHRG---IEMDKMNSH 306
 QY 296 ---FLDGNLPLEGKEVD---ESRRIRIKLKDOKHPKEDDOLVEMANYALSHQ 348
 DB 307 VENLDGALV-LEVDERDQDEEARREMARILKELKOKHPKEDDOLVEMANYALSHQ 365
 QY 349 KSRAFYRIQATRMATGAGNLIKHAADQAKKASSMEVHDEPE-DFISKVFPDPCSYOC 407
 DB 366 KSRAFYRIQATRMATGAGNLIKHAADQAKKASSMEVHDEPE-DFISKVFPDPCSYOC 425
 QY 408 LENGAVLLTVRKGGDMKTMVDTKTEDGSANAGADYEFTETVYLKPEOTKESVYG 467
 DB 426 LENGAVLLTVRKGGDMKTMVDTKTEDGSANAGADYEFTETVYLKPEOTKESVYG 485
 QY 468 IIDDIFEEDHEFVRLSNVRIEEOPEEGMPAIFNSLPLPRVILASPCVATYTIIDD 527
 DB 486 IIDDIFEEDHEFVRLSNVRIEEOPEEGMPAIFNSLPLPRVILASPCVATYTIIDD 542
 QY 528 HAGIFTECDTIHVESIGVAVKLTSGARGTVIVPFTVEGTAGGGEDEFTDYGEL 587
 DB 543 HAGIFTECDTIHVESIGVAVKLTSGARGTVIVPFTVEGTAGGGEDEFTDYGEL 602
 QY 588 EFKNDE 593
 DB 603 EFKNDE 608

RESULT 12

ABB61721

AC ABB61721;

DT 26-MAR-2002 (first entry)

DE Drosophila melanogaster polypeptide SEQ ID NO 11955.

XX Drosophila; developmental biology; cell signalling; insecticide;
 KW pharmaceutical.

XX Drosophila melanogaster.

XX WO200171042-A2.

XX 27-SEP-2001.

XX 23-MAR-2001; 2001WO-US09231.

XX 23-MAR-2000; 2000US-191637P.

XX 11-JUL-2000; 2000US-0614150.

XX (PEKE) PE CORP NY.

XX Venter JC, Adams M, Li PWD, Myers EW,

XX WPI; 2001-656860/75.

XX N-PSDB; ABL05824.

XX New isolated nucleic acid detection reagent for detecting 1000 or more
 PT genes from Drosophila and for elucidating cell signalling and cell-cell
 PT interactions -

PS Disclosure; SEQ ID NO 11955; 21pp + Sequence Listing; English.

CC The invention relates to an isolated nucleic acid detection reagent
 CC capable of detecting 1000 or more genes from Drosophila. The invention is
 CC useful in developmental biology and in elucidating cell signalling and
 CC cell-cell interactions in higher eukaryotes for the development of
 CC insecticides, therapeutics and pharmaceutical drugs. The invention
 CC discloses genomic DNA sequences (AB16176-AB130511), expressed DNA
 CC sequences (AB101840-AB116175) and the encoded proteins
 CC (ABBS7737-ABBS72072).

CC The sequence data for this patent did not form part of the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.

XX Sequence 950 AA;

Query Match 44.4%; Score 2130.5; DB 22; Length 950;

Best Local Similarity 49.2%; Pred. No. 2.8e-198;

Matches 436; Conservative 160; Mismatches 235; Indels 56; Gaps 16;

QY 53 DCKRGVLLPIWYR-NSLDGKIRARIVYFVALIYMLGVSIIADREMAIEVITTSQRE 111
 DB 99 ECSEGLVLPMPORNTSVGDRVGFVYLLVFGVSIADREMAIEVITTSQRE 158
 QY 112 VTIRKPNGETSTTIRVWNETVSNLTLMALGSSAPETLSIEVCGHGFITGDIQSPSTIV 171
 DB 159 VYVAGPNTKQVMIRINENAVNLTLMALGSSAPETLSIEVCGHGFITGDIQSPSTIV 218
 QY 172 GSAFNMFIIGICVYVDPDETRKIKHLRVFTTAWSIFAYIMLYMILAVFSPGVQV 231
 DB 219 GSAFNMFIIGICVYVDPDETRKIKHLRVFTTAWSIFAYIMLYMILAVFSPGVQV 278
 QY 232 WEGILLTFEPFVVCVLAMVADKRLFFYMKKRTDKHKGIIETESDHRG---IEMDKMNSH 290
 DB 279 WEGILLTFEPFVVCVLAMVADKRLFFYMKKRTDKHKGIIETESDHRG---IEMDKMNSH 334
 QY 291 ----MNSHFLDGNLPLEGKEVDSESRIRIKLKDOKHPKEDDOLVEMANYALSHQ 345
 DB 335 GPKOPWTS--ARGN----DAEAFDEARREYITLTTELKOKYPADLEQLEMAQEOVLA 388
 QY 346 HOKSRAFYRIQATRMATGAGNLIKHAADQAKKASSMEVHDEPE-DFISKVFPDPCSYOC 399
 DB 389 RSSKSRAFYRIQATRMATGAGNLIKHAADQAKKASSMEVHDEPE-DFISKVFPDPCSYOC 445
 QY 400 FDCPSYOCLENGAVLLTVRKGGDMKTMVDTKTEDGSANAGADYEFTETVYLKPEOTKESVYG 459
 DB 446 FEPGHTVWENCGEFEEVAVVR-GDISTVASVEYETODGTASAGTDVFGKGLSLSPPGV 504

OY 460 TOKESVGIIDDDIFEDEHFFRLSNVRLNEEQPEEGMPAIFNSLPLBRAVLASPCA 519
DB 505 DQRRRIEYIDDDVEEDCEFYIRLEN-----PSEGVK-----LAVPMIA 544
OY 520 TWTIIDDHAGJFTEPCDTIHVESISIGVMEVKVLTSGARGVYVFPFTEGTAKGGED 579
DB 545 TWTIIDDHAGJFATFDSVFELTESYGRFELKMRSGARGVYVYWTENDPAT-ESKD 603
OY 580 FEDTYGELEFKDEYVKTIRKVIDEEXERQDNFIALGEPKW-----MENGISVYDR 634
DB 604 YEGARSELVEFNNESEKFDLPLEESSYKEDVSFKVHIGEPRLADDELAIAKIKEV-EK 662
OY 635 KILMESEAKRIAMCKPVLGEHPKLEVIIEESYEKKTVDKIKKTNALVYGTSMRD 694
DB 663 KPVQDITELDRILLKSPKNGELTAYVRIRESQERKATVDKLVAAANSAVLGTSMKR 722
OY 695 QPMEAITVSAGD-EDEDESGEERLPCSPDYVNHFLTFVEMKVLACVPPTXCHGACPA 753
DB 723 QPKDALVIPADESEFENDDEEYVSCFSYSHFVCLRWKVLFAVVPPTDIOGCVTVY 782
OY 754 VSILITGMLTAIIGDLASHFGCTIGLSDSVTAVVAVFSTVDPTEFASKAALQDYYADA 813
DB 783 VSIFVIGVITAIIGDAASYFGCALNKIDSVTALFVALGTSIPDTPASMIAAKHDEGADN 842
OY 814 SIGNVTSNAVNVFLGIGLAMSVAATYMALQGEHFVSAAGTLAFSVTLTIFAFVCISVL 873
DB 843 CIGNVTSNAVNVFLGIGLAMSVAATYMALQGEHFVSAAGTLAFSVTLTIFAFVCISVL 902
OY 874 LYRR-RPHLGELGPRGCKLATTWLVSIMLXYLFATLEAVCYIK 919
DB 903 MFRRMHKGIGALGPRKYSKYSIAALVLEWVYVICILEANDVIR 949

RESULT 13

ABG26781
ID ABG26781 standard; Protein; 394 AA.

AC ABG26781;

DT 18-FEB-2002 (first entry)

DE Novel human diagnostic protein #26772.

Human; chromosome mapping; gene mapping; gene therapy; forensic;
food supplement; medical imaging; diagnostic; genetic disorder.

Human sapiens.

MO200175067-AA2.

11-OCT-2001.

30-MAR-2001; 2001WO-US08631.

31-MAR-2000; 2000US-0540217.

23-AUG-2000; 2000US-0649167.

(HYSE-) HYSEQ INC.

Dmanac RT, Liu C, Tang YT;

WPI: 2001-639362/73.

N-PSDB; AAS90968.

New isolated polynucleotide and encoded polypeptides, useful in
diagnostics, forensics, gene mapping, identification of mutations
responsible for genetic disorders or other traits and to assess
biodiversity

Claim 20; SEQ ID NO 57140; 103bp; English.

The invention relates to isolated polynucleotide (1) and

CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (II) is useful in gene therapy techniques
CC to restore normal activity of (I) or to treat disease states involving
CC (II). (II) is useful for generating antibodies against it, detecting or
CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. ABG00010-ABG3037 represent novel human
CC diagnostic amino acid sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC at ftp.wipo.int/pub/published_pct_sequences.

Sequence 394 AA:

Query Match 33.5%; Score 1607; DB 22; Length 394;
Best Local Similarity 83.0%; Pred. No. 9, 2e-148;
Matches 313; Conservative 19; Mismatches 31; Indels 14; Gaps 3;

OY 559 RGTIVVPFTEGTAKGCG---EDFEDTYGELEFKN-----DETIVKTRKIVDE 605
DB 18 RODAVMPNRRKNKKNYKKNPAKRCASESSHGKVRSSSTCVQLPYKEALKTIRKVIDD 77
OY 606 EYERQDNFIALGEPKMERGI-SDVTDKRLMESEAKRIAMCKPVLGEHPKLEVI 664
DB 78 EAYEKKNKVFTEEMGPRMVDMSFQKDYTDKRLMESEAKRIAMCKPVLGEHPKLEVI 137
OY 665 EESYEKKTVDKLIKKTNLALVYGTSMRDQFMEAITVSAAGDEDESGEERLPCSPDY 724
DB 138 EESYEKKTVDKLIKKTNLALVYGTSMRDQFMEAITVSAAGDEDESGEERLPCSPDY 197
OY 725 VMHFLTFVEMKVLACVPPTXCHGMACFAYSLIIGMLTAIIGDLASHFGCTIGLSDSV 784
DB 198 VMHFLTFVEMKVLACVPPTXCHGMACFAYSLIIGMLTAIIGDLASHFGCTIGLSDSV 257
OY 785 AVYFAFGTSVPTPFASKAALQDYYADASIGNVTSNANVFLGIGLAMSVAATYMALQ 844
DB 258 AVYFAFGTSVPTPFASKAALQDYYADASIGNVTSNANVFLGIGLAMSVAATYMALQ 317
OY 845 GQEFHVSAGTLAFSVTLTIFAFVCISVLLYRRRPHLGELGPRGCKLATTWLFPVSLML 904
DB 318 GQEFHVSAGTLAFSVTLTIFAFVCISVLLYRRRPHLGELGPRGCKLATTWLFPVSLML 377
OY 905 LYTLFATLEAVCYIKGF 921
DB 378 LYTLFATLEAVCYIKGF 394

RESULT 14

AAO05893
ID AAO05893 standard; Protein; 120 AA.

AC AAO05893;

DT 06-NOV-2001 (first entry)

Human polypeptide SEQ ID NO 19785.

Human; cytokine; cell proliferation; cell differentiation; gene therapy;
vaccine; peptide therapy; stem cell growth factor; haematopoiesis;
tissue growth factor; immunomodulatory; cancer; leukaemia;
nervous system disorders; arthritis; inflammation.

Homo sapiens.

OS

PN W0200164835-A2.
XX
PD 07-SEP-2001.
XX
PE 26-FEB-2001; 2001WO-US04927.
XX
PR 28-FEB-2000; 2000US-0515126.
XX 18-MAY-2000; 2000US-0577409.
XX
PA (HYSE-) HYSEQ INC.
XX
PI Tang YT, Liu C, Drmanac RT;
XX WPI; 2001-514838/56.
DR N-PSDB; AA185824.
XX
PT Isolated nucleic acids and polypeptides, useful for preventing
PT diagnosing and treating e.g. leukemia, inflammation and immune
PT disorders -
XX
PS Claim 20: SEQ ID NO 19785; 1399pp + Sequence Listing; English.

CC The invention relates to human polynucleotides (AA179941-AA193841) and
CC the encoded proteins (AA000010-AA013910) that exhibit activity elating to
CC cytokine, cell proliferation or cell differentiation or which may induce
CC production of other cytokines in other cell populations. The
CC polynucleotides and polypeptides are useful in gene therapy, vaccines or
CC peptide therapy. The polypeptides have various cytokine-like activities,
CC e.g. stem cell growth factor activity, haematopoiesis regulating
CC activity, tissue growth factor activity, immunomodulatory activity and
CC activin/inhibin activity and may be useful in the diagnosis and/or
CC treatment of cancer, leukemia, nervous system disorders, arthritis and
CC inflammation.
CC Note: The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at [ftp://www.int/pub/published_pct_sequences](http://www.int/pub/published_pct_sequences).
XX
SQ Sequence 120 AA;
XX

Query Match 12.5%; Score 599; DB 22; Length 120;
Best Local Similarity 100.0%; Pred. No. 4,2e-50;
Matches 120; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 757 LIIGMLTAIIGDLASHFECTIGLDSYAVVFAFGTSVPTFASKAALQDVYADASIG 816
DB 1 LIIGMLTAIIGDLASHFECTIGLDSYAVVFAFGTSVPTFASKAALQDVYADASIG 60
QY 817 NVTSSNAVNVFLGIGLMSVAIYAWALOGDFHVSAGTLAFSVTLFTIFAFVCISVLLXR 876
DB 61 NVTSSNAVNVFLGIGLMSVAIYAWALOGDFHVSAGTLAFSVTLFTIFAFVCISVLLXR 120

RESULT 15
AA158044
ID AA158044 standard; Protein; 539 AA.
XX
AC AA158044;
XX
DT 14-MAR-2000 (first entry)
XX
DE Arabidopsis thaliana 11 transmembrane domain transporter MXH.
XX
KM 11 transmembrane domain transporter; MXH; metal cation-proton exchanger;
KM zinc; magnesium; stress tolerance; transgenic plant.
XX
OS Arabidopsis thaliana.
XX
PN W09961616-A2.
XX
PD 02-DEC-1999.
XX
PE 25-MAY-1999; 99WO-IL00277.
XX

PR 26-MAY-1999; 98IL-0124653.
XX
PA (YEDA) YEDA RES & DEV CO LTD.
PA (FLAN-) FLANDER INTERUNIVERSITY INST BIOTECHNOLO.
PA (TEXA) UNIV TEXAS SYSTEM.
XX
PI Gallili G, Shaul O, Inze D, Van Montagu M, Hilgemann DW;
XX WPI; 2000-062714/05.
DR N-PSDB; AA247475, AA247476.
XX
PT New isolated DNA encoding a metal cation-proton exchanger from plants,
PT used to produce transgenic plants with improved stress tolerance
XX
PS Claim 3; Fig 1; 52pp; English.

CC This is the Arabidopsis thaliana 11 transmembrane domain transporter
CC designated MXH protein sequence. The MXH protein is a member of the 11-12
CC transmembrane domain transporter family and has Mg²⁺/H⁺ or Zn²⁺/H⁺
CC exchange activity. In plants Mg²⁺ and Zn²⁺ are essential for the
CC integrity of ribosomes and cellular function require a fine balance of
CC various ions including magnesium and zinc. The polypeptide and nucleotide
CC sequences of MXH can be used to transform plant cells and produce
CC transgenic plants. The transformed plants have improved tolerance to
CC stress conditions such as drought, temperature, mineral excess or
CC deficiency and high salinity. Other improvements include better growth
CC on calcareous soils, possibly also improved uptake of important metals
CC where these have limited availability in the soil.
XX
SQ Sequence 539 AA;
XX

Query Match 12.1%; Score 579; DB 21; Length 539;
Best Local Similarity 22.3%; Pred. No. 5e-47;
Matches 191; Conservative 110; Mismatches 199; Indels 356; Gaps 22;

QY 64 YPENPSGDKIARIYIVFALITYFLGVSITADFFMASIEYITSOEDEVITKPKNGEST 123
DB 29 FPGENTLSDDL-RGVLYFLGLAYCFIGLSAITARFFKSMENVVHSHKRYVITPITRAEV 87
QY 124 TTI-RVNNETYSNLTLMAGSSAPEILLSTIEV---CGHGFIADLPSTIVGSAANMF 179
DB 88 ITTKVNNFTIADISLAFGTSFPOISLATIDAIRNNGERY-AGGLPRTIVGSAARDLF 146
QY 180 IIGICVYVYPDGETRKIKHLRVEFITAAVSIFAYIMLYMLAFSPGVVOWEGELTLF 239
DB 147 PIHAVCVVPRAGLKKIKSIDGLVGLVLEWVSFWAYIMLYITLEWSPNVITLVEALLTVL 206
QY 240 FFPYCVLLAVVADKRLLFYKYMKKYRTDKHGIITETEGDHPKIGMDGKMNSHFLDG 299
DB 207 QYGLLVHAAVADKR--WPLTS-----LPMRSGDRPE----- 236
QY 300 NLVPLEGKEWDESREMIRILDKQKHPEKDLQLEVMANYVALSHQKSRARFRIQAT 359
DB 237 EWVP---EIDTSK-----DDNDND----- 253
QY 360 RMTAGAGNIILKHAEOAKKASSMSEVHTDEPEDFISKVEFPDPCSYOCLENCAVLLTVV 419
DB 254 -----VHDVYSDAADAIV----- 266
QY 420 RKGDMKSTMYVDYKTEGDSANAGADYETEGTVVLKPGTQKEFSVGIIDDDIFEEDEH 479
DB 267 -----ESGSRN-----IVD----- 275
QY 480 FFWLNSVNRIEBQPEGMPAIFNSLPRLRAVLASPCAVTITLDDDHAGITFEEDCTI 539
DB 276 -----II-----SI 279
QY 540 HVSESIGMEVKVLTSGARGTVIVPRVEGTAKGGEDEFTYGELEFRKNDETVTIR 599
DB 280 H-----SANNDCITHTYADTRP-----DSATKKK 306
QY 600 VKIVDEEYERQENFTALGEPKMERGISDVTDRKLIMEEPAKRIAEKGPVLEGHPR 659

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Db 307 AK----- 308
Qy 660 LEVIESEYFKTYDKLIKKTNLALVGHSHMRDQMEATVYSAGDEDESEGERLP 719
Db 309 -----NSTVFDI-----WKHOFVDATVLETSESKVD-STYLRJA 342
Qy 720 SCFDYVMHFLTVFWKVLFCVPTPEYCHGWACPAVSLIIGMLTAIIGDLASHFGCTIGL 779
Db 343 KSF---WHLLAPWKLLFAFVPPCNTAHGMIATCSLFLTSGVAFYVTRFDLSCVTGI 399
Qy 780 KDSVAVVFAVGTSVPTFASKAALQDVYADASTGNVTSNANVFLGIGLANSVAAT 839
Db 400 NPVIAFTALASGTSWPDVASKIAEROLTADSAIANITCSNSNIVYIGVPMILINTV 459
Qy 840 --YWALOGGEFHSAGTLAFSVTLFTIIFAVCISVLLYRRRPHLGELGPRGCKLATW 897
Db 460 YNFFAYREPLYTENAKGLSPSLIFPATSVGCIVLVLRRL-IIGAE LGGPRLMAMLTSA 518
Qy 898 LFWSLMLLYILFATLE 913
Db 519 YFMMLMVVFVLSLTK 534

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